Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) A method of delivering a therapeutic dose of a bioactive agent to the pulmonary system of a subject, in a single, breath-activated step, comprising:

administering particles comprising a bioactive agent, from a receptacle having a mass of particles, to a subject's respiratory tract, wherein:

i) the particles administered to the subject's respiratory tract have a tap density of less than 0.4 g/cm^3;

ii) ~~[a)~~ at least 50% of the particles have a fine particle fraction ~~less than 4.0 μm; and/or~~

~~b)]~~ at least 75% of the particles have a fine particle fraction less than 6.8 μm; and

iii) at least about 50% of the mass of particles stored in the receptacle is delivered to the pulmonary system of the subject.

2. (original) The method of Claim 1 wherein the particles have a tap density of less than about 0.1 g/cm^3.

3. (original) The method of Claim 1 wherein the particles have a geometric diameter greater than about 5 μm.

4. (original) The method of Claim 1 wherein the receptacle has a volume of at least about 0.37 cm^3.

5. (previously amended) The method of Claim 1 wherein the receptacle has a volume of at least about 0.48 cm^3.

6. (previously amended) The method of Claim 1 wherein the receptacle has a volume of at least about 0.67 cm^3.

7. (previously amended) The method of Claim 1 wherein the receptacle has a volume of at least about 0.95 cm^3.

8. (original) The method of Claim 1 wherein delivery is primarily to the deep lung.

9. (original) The method of Claim 1 wherein delivery is primarily to the central airways.

10. (original) The method of Claim 1 wherein the bioactive agent is albuterol sulfate.

11. (original) The method of Claim 1 wherein the bioactive agent is insulin.

12. (original) The method of Claim 1 wherein the bioactive agent is growth hormone.

13. (original) The method of Claim 1 wherein the bioactive agent is fluticasone.

14. (original) The method of claim 1 wherein the bioactive agent is salmeterol.

15. (original) The method of Claim 1 wherein the bioactive agent is a hydrophobic drug.

16. (original) The method of Claim 1 wherein the bioactive agent is a hydrophilic drug.

17. (original) The method of Claim 1 wherein the bioactive agent is a monoclonal antibody.

18. (original) The method of Claim 1 wherein the particles are in the form of a dry powder.

19. (original) The method of Claim 1 wherein administration to the respiratory tract is by a dry powder inhaler.

20. (currently amended) A method of delivering a therapeutic dose of a bioactive agent to the pulmonary system of a subject, in a single breath, comprising:

administering dry powder particles comprising a bioactive agent, from a receptacle having a mass of particles, to a subject's respiratory tract in a single breath,

wherein:

i) the particles have a tap density less than about 0.4 g/cm^3; [and]

ii) at least about 5 milligrams of the bioactive agent is delivered to the pulmonary system of the subject[.] and

iii) at least 75% of the particles have a fine particle fraction less than 6.8 μm.

21. (original) The method of Claim 20 wherein the particles have a tap density of less than about 0.1 g/cm^3.

22. (original) The method of Claim 20 wherein the particles have a geometric diameter greater than about 5 μm.

23. (original) The method of Claim 20 wherein the receptacle has a volume of at least about 0.37 cm^3.

24. (previously amended) The method of Claim 20 wherein the receptacle has a volume of at least about 0.48 cm^3.

25. (previously amended) The method of Claim 20 wherein the receptacle has a volume of at least about 0.67 cm^3.

26. (previously amended) The method of Claim 20 wherein the receptacle has a volume of at least about 0.95 cm^3.

27. (original) The method of Claim 20 wherein the particles deliver at least 15 milligrams of the bioactive agent.

28. (original) The method of Claim 20 wherein the particles deliver at least 20 milligrams of the bioactive agent.

29. (original) The method of Claim 20 wherein the particles deliver at least 30 milligrams of the bioactive agent.

30. (original) The method of Claim 20 wherein the particles deliver at least 35 milligrams of the bioactive agent.

31. (original) The method of Claim 20 wherein the particles deliver at least 50 milligrams of the bioactive agent.

32. (original) The method of Claim 20 wherein delivery is primarily to the deep lung.

33. (original) The method of Claim 20 wherein delivery is primarily to the central airways.

34. (original) The method of Claim 20 wherein the bioactive agent is albuterol sulfate.

35. (original) The method of Claim 20 wherein the bioactive agent is insulin

36. (original) The method of Claim 20 wherein the bioactive agent is growth hormone.

37. (original) The method of Claim 20 wherein the bioactive agent is ipratropium bromide.

38. (original) The method of Claim 20 wherein the bioactive agent is fluticasone.

39. (original) The method of claim 20 wherein the bioactive agent is salmeterol.

40. (original) The method of Claim 20 wherein the bioactive agent is a hydrophobic drug.

41. (original) The method of Claim 20 wherein the bioactive agent is a hydrophilic drug.

42. (original) The method of Claim 20 wherein the bioactive agent is a monoclonal antibody.

43. (original) The method of Claim 20 wherein the particles are in the form of a dry powder.

44. (original) The method of Claim 20 wherein administration to the respiratory tract is by a dry powder inhaler.

45. (previously presented) The method of Claim 1 wherein at least 50% of the particles have a fine particle fraction less than 4.0 μm.

46. Canceled

47. (previously presented) The method of Claim 20 wherein at least 50% of the particles have a fine particle fraction less than 4.0 μm.

48. Canceled

49. (previously presented The method of Claim 1 wherein said particles are spray dried particles.

50. (previously presented) The method of Claim 20 wherein said particles are spray dried particles.

51. (previously presented) The method of Claim 20 wherein the particles deliver at least 10 milligrams of the bioactive agent.